Appl. No.

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AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

- 1. (Currently amended) A recombinant plasmid vector which comprises:
 - a kanamycin resistance gene;
 - a promoter;
 - a nucleotide sequence coding for an endoxylanase signal sequence;
- a nucleotide sequence coding for an oligopeptide consisting of 13 amino acids, wherein-including 6 of the 13 amino acids are consecutive histidine residues; and,
 - a human granulocyte colony stimulating factor (hG-CSF) gene.
- 2. (Currently amended) The recombinant plasmid vector of claim 1, wherein the nucleotide sequence codes coding for an the oligopeptide which comprises an amino acid sequence of isoleucine glutamic acid glycine arginine (Ile-Glu-Gly-Arg; comprises a nucleic acid encoding SEQ ID NO: 28) within the oligopeptide.
- 3. (Currently amended) A recombinant plasmid vector, pTHKCSFmII represented in Figure 13 which comprises:
 - a kanamycin resistance gene;
 - a Trc promoter;
 - a nucleotide sequence coding for an <u>a Bacillus sp.</u> endoxylanase signal sequence derived from Bacillus sp.;
 - a nucleotide sequence coding for the oligopeptide of SEQ ID NO: 1; and
 - a modified gene coding for a human granulocyte colony stimulating factor (hG-CSF) lacking its native signal sequence.
- 4. (Currently amended) A microorganism, E. coli transformed with the plasmid vector, pTHKCSFmII of claim 3.
- 5. (Currently amended) The E. coli microorganism-of claim 4, wherein the E. coli is selected from the group consisting of E. coli XL1-Blue, E. coli MC4100, E. coli BL21 (DE3), E. coli HB101 and E. coli W3110.

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6. (Currently amended) E. coli MC4100/pTHKCSFmII, deposited as (KCTC 0754BP), wherein said E. coli is transformed with the plasmid vector, pTHKCSFmII of claim 3.

7. (Original) A process for preparing a human granulocyte colony stimulating factor, which comprises the steps of:

culturing *E. coli* transformed with the plasmid vector of claim 1 to obtain a human granulocyte colony stimulating factor fusion protein; and,

treating the human granulocyte colony stimulating factor fusion protein with a protease to obtain a human granulocyte colony stimulating factor.

- 8. (Currently amended) The process for preparing a human granulocyte colony stimulating factor of claim 7, wherein the plasmid vector of claim 1 is pTHKCSFmII.
- 9. (Currently amended) The process for preparing a human granulocyte colony stimulating factor of claim 7, wherein the human granulocyte colony stimulating factor fusion protein is <u>isolated from the protein pool</u> obtained from the culture <u>using a Ni-column</u>.
- 10. (Original) The process for preparing a human granulocyte colony stimulating factor of claim 7, wherein the protease is Factor Xa.
- 11. (Previously added) The recombinant plasmid vector of Claim 3, wherein said vector comprises the nucleotide sequence of SEQ ID NO: 26.
- 12. (Currently amended) The recombinant plasmid vector of Claim 3, wherein said modified gene comprises nucleotides 88 to 610 of the <u>nucleotide sequence of SEQ</u> ID NO: 18 and encodes the hG-CSF amino acid sequence of SEQ ID NO: 19.
- 13. (Previously added) The recombinant plasmid vector of Claim 3, wherein said nucleotide sequence coding for said endoxylanase signal sequence comprises nucleotides 1-84 of SEQ ID NO: 26.